PATENT COOPERATION TREATY

| From the INTERNATIONAL SEARCHING AUTHORITY | | • | | | |
|---|--|--|---|--|--|
| To: IVOR R. ELRIFI MINTZ, LEVIN, COHN, FERRIS GLOVSKY AND POPEO PC | | PCT | | | |
| ONE FINANCIAL CENTER BOSTON, MA 02111 | | WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY | | | |
| | | | (PCT Rule 43bis.1) | | |
| | | Date of mailing (day/month/year) | 0 1 NOV 2005 | | |
| Applicant's or agent's file reference | | FOR FURTHER ACTION See paragraph 2 below | | | |
| 24028-015-06 | | | | | |
| International application No. International filing date | | | Priority date (day/month/year) | | |
| PCT/US05/10744 29 Mar International Patent Classification (IPC) or both n | ch 2005 (29.03.2 | tion and IPC | 29 March 2004 (29.03.2004) | | |
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| IPC(7): C12N 15/00; C12P 19/34 and US Cl.: 43 Applicant | 35/440, 91.1 | | | | |
| TABATADZE ET AL | | | · | | |
| TABATADZE ET AL | | | | | |
| 1. This opinion contains indications relating to the following items: | | | | | |
| Box No. I Basis of the opinion | | | | | |
| Box No. II Priority | | | | | |
| Box No. III Non-establishment of | Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability | | | | |
| Box No. IV Lack of unity of invention | | | | | |
| Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement | | | | | |
| Box No. VI Certain documents co | Box No. VI Certain documents cited | | | | |
| Box No. VII Certain defects in the | Box No. VII Certain defects in the international application | | | | |
| Box No. VIII Certain observations on the international application | | | | | |
| 2. FURTHER ACTION | | | | | |
| If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered. | | | | | |
| If this opinion is, as provided above, conside IPEA a written reply together, where appromailing of Form PCT/ISA/220 or before the For further options, see Form PCT/ISA/220. | opriate, with am expiration of 22 | endments, before the | ne expiration of 3 months from the date of | | |
| 3. For further details, see notes to Form PCT/IS | | | | | |
| Name and mailing address of the ISA/ US Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 | Date of comple opinion 19 September 2 | tion of this 2005 (19.09.2005) | Authorized officer Tara L. Garvey Telephone No. (571) 272-0507 | | |

Facsimile No. (703) 305-3230
Form PCT/ISA/237 (cover sheet) (April 2005)

WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.

PCT/US05/10744

| Box No. I Basis of this opinion |
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| 1. With regard to the language, this opinion has been established on the basis of: |
| the international application in the language in which it was filed |
| a translation of the international application into, which is the language of a translation furnished for the purposes of international search (Rules 12.3(a) and 23.1(b)). |
| 2. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of: |
| a. type of material |
| a sequence listing |
| table(s) related to the sequence listing |
| b. format of material |
| On paper |
| in electronic form |
| c. time of filing/furnishing |
| contained in the international application as filed. |
| filed together with the international application in electronic form. |
| furnished subsequently to this Authority for the purposes of search. |
| |
| In addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished. |
| 4. Additional comments: |
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WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/US05/10744

Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

| 1. Statement | | |
|-------------------------------|--------------------------------|-----|
| Novelty (N) | Claims 26-40 | YES |
| • | Claims <u>1-5, 7-19, 21-25</u> | NO |
| Inventive step (IS) | Claims <u>26-40</u> | YES |
| | Claims 1-25 | |
| Industrial applicability (IA) | Claims 1-40 | YES |
| | Claims NONE | NO |

2. Citations and explanations:

Claims 1-5, 7-19 and 21-25 lack novelty under PCT Article 33(2) as being anticipated by anticipated by Pederson et al. Pederson et al teaches a method of site-directed alteration of an RNA molecule using an oligonucleotide that is flanked by nucleotide sequences unable to activate RNase H and a second oligonucleotide that is unmodified and hybridized to the moficied oligonucleotides and altering the RNA in the presence of Rnase. Furthermore, the alteration can be excision or excision and addition of nucleotides, the oligonucleotides can be modified with phosphorothioates, the flanking sequences can be modifies with methyl phosphonates. The method may be used to correst defects in an individual suffering from cystic fibrosis (abstract, column 1, lines 59-67, columns 2-6 and columns 11-12).

Claims 6 and 20 lack an inventive step under PCT Article 33(3) as being obvious over Pederson et al in view of Kole et al. Pederson et al does not specifically teach modifying the oligonucleotide by adding a 2-O-methyl moiety. Kole et al teaches modfying oligonucleotides with a 2-O-methyl moiety for alteration of RNA molecules (column 7, lines 44-67 and column 8, lines 1-7). In view of this teaching, it would have been obvious to one of ordinary skill in the art to modify the oligonucleotides with a 2-O-methyl moiety as described by Kole et al.

Claims 26-40 meet the criteria set out in PCT Article 33(2)-(3), because the prior art does not teach or fairly suggest a method of repairing a gene by contacting an RNA molecule with a hybrid DNA/RNA oligonucleotide complex or an RNA oligonucleotide complex comprising SEQ ID NO:1 and SEQ ID NO:2.

Claims 1-40 meet the criteria set out in PCT Article 33(4), and thus have industrial applicability because the subject matter claimed can be made or used in industry.